Amendments to the Claims

This listing of the claims will replace all prior versions and listing of claims in this application.

Listing of Claims

1. (Currently Amended) A liquid pharmaceutical preparation comprising pradofloxacin bound to an ion exchange resin, characterized in that the loaded ion exchange resin is dispersed in a carrier medium comprising water and one or more pseudoplastic gel formers**, wherein the pseudoplastic gel former is selected from the group consisting of polyacrylic acid, xanthan, microcrystalline cellulose, cellulose ether, bentonite, and a mixture thereof**.
2. (Canceled).
3. (Currently Amended) The pharmaceutical preparation according to claim 1, wherein the ~~the~~ ion exchange resin is an acidic ion exchanger.
4. -10. Canceled.
11. (Previously presented) The pharmaceutical preparation according to claim 1, wherein the pseudoplastic gel former is xanthan.
12. (Previously presented) The pharmaceutical preparation according to claim 1, wherein pradofloxacin loading on the ion exchange is from about 1% to about 50% by weight.
13. (Previously presented) The pharmaceutical preparation according to claim 1, wherein pradofloxacin loading on the ion exchange is from about 5% to about 30% by weight.
14. (Previously presented) The pharmaceutical preparation according to claim 1, wherein the carrier medium is from about 10 to about 98 % by weight of the total preparation.
15. (Previously presented) The pharmaceutical preparation according to claim 1, wherein the carrier medium is from about 20% to about 90% by weight of the total preparation.

16. (Previously presented) The pharmaceutical preparation according to claim 1, wherein the yield point of the preparation is between 0 and 100 Pa.
17. (Previously presented) The pharmaceutical preparation according to claim 1, wherein the yield point of the preparation is between 5 and 50 Pa.
18. (Previously presented) The pharmaceutical preparation according to claims 1, wherein the viscosity at 300 s^{-1} of the preparation is between 10 and 1000 mPa*s
19. (Previously presented) The pharmaceutical preparation according to claims 1, wherein the viscosity at 300 s^{-1} of the preparation is between 50 and 500 mPa*s.